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MN Relay Service for Hearing Impaired 800.627.3529

## **ADVERSE REACTION REPORT**

Minnesota Rules 3100.3600 requires that you file this report for any incident that arises from the administration of nitrous oxide inhalation analgesia or of a pharmacological agent for the purpose of general anesthesia, conscious sedation, local anesthesia, analgesia, or anxiolysis that results in a serious or unusual outcome that produces a temporary or permanent physiological injury, harm, or other detrimental effect to one or more of a patient's body system(s). It is <u>NOT</u> necessary to report incidents such as nausea, a single episode of emesis, or mild allergic reaction. This report shall be submitted to the Board of Dentistry within ten days of the incident. You may duplicate this form.

LICENSEE INFORMATION					
Name (please print):	License Number:				
Addroop					
Address:					
City:	State:			Zip	
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I. REACTION INFORMATION					
PATIENT ID/INITIALS (In Confidence)	AGE SEX (YRS)	REACTION ONSET			
	(1K3)	МО	DA	YR	CHECK ALL APPROPRIATE:
DESCRIBE REACTION(S)					<b>—</b>
					PATIENT DIED
					REACTION TREATED WITH RX DRUG
					RESULTED IN TREATMENT BY PHYSICIAN AND/OR HOSPITALIZATION
RELEVANT TESTS/LABORATORY DATA				RESULTED IN PERMANENT DISABILITY	
					NONE OF THE ABOVE
11.	SUSPECT DRUG	G(S) INFO	RMATION		
SUSPECT DRUG(S) (Indicate manufacturer and lot # for vaccines/biologics)				DID REACTION ABATE AFTER STOPPING DRUG?	
DOSE	ROUTE OF ADM	ROUTE OF ADMINISTRATION			YES NO NA
INDICATION(S) FOR USE					DID REACTION REAPPEAR AFTER REINTRODUCTION?
DATES OF ADMINISTRATION (From/To)	DURATION OF A	DURATION OF ADMINISTRATION			YES NO NA
III.	CONCOMITANT D			RY	<u> </u>
CONCOMITANT DRUGS AND DATES OF ADMINISTR	ATION (Exclude those used to	treat reaction)			
OTHER RELEVANT HISTORY (e.g., diagnoses, allergie	es, pregnancy with LMP, etc.)				
IV.	eic	NATURE			
IV.	310	NATURE			
SIGNED:	DATE:				